Orthophosphate by SEAL AQ2 Discrete Analyzer SEAL Method EPA-118-A Rev. 3						Page 1 of 2
Facility Name:	VELAP ID					
Assessor Name:Analyst Name:	Inspection Date					
Relevant Aspect of Standards	Method Reference	Y	N	N/A	Comments	
Records Examined: SOP Number/ Revision/ Date				A	nalyst:	
Sample ID: Date of Sample Prepare	ration:		Da	ate of A	Analysis:	
Is the linear calibration range determined initially, and does it contain a minimum of a blank and three standards?	Method Supplement 1, Rev. 2 (MS) 3.2.1					
Is linearity reestablished if any verification data exceeds initial calibration values by ±10%?	MS 3.2.1					
Is a laboratory control sample analyzed with every batch, and is recovery within ±10% of the stated value?	мs 3.4.3					
Are method detection limits established?	мs 3.4.3					
Is at least one method blank carried through all the procedural steps with each batch?	мs 3.4.1.1					
Is the initial calibration verified using a second source or certified standard other than the quality control sample?	мѕ 4.4					
Is the calibration verified using a calibration standard after every ten samples or every analytical batch?	MS 4.5					
Is a minimum of 10% of all samples spiked with the stock standard?	MS 3.3.1					
If matrix interference is present, are results not reported for regulatory compliance purposes?	MS 3.3.1.4.1					
For compliance monitoring, is the concentration of the matrix spike at the regulatory limit OR 1 to 5 times higher than the background concentration of the sample?	MS 3.3.1.1.1					
For dissolved orthophosphate for EPA's wastewater compliance monitoring programs, are samples filtered immediately after collection, cooled to 4°C, and analyzed within 48 hours? (No acid is used.)	8.3					
Notes/Comments:						

Relevant Aspect of Standards    Method Reference   Y   N   N/A   Comments	Orthophosphate by SEAL AQ2 Discrete Analyzer SEAL Method EPA-118-A Rev. 3 Page 2 of 2						
monitoring programs, are samples NOT filtered, but cooled to 4°C and analyzed within 48 hours? (No acid is used.)  Are the AQ2 test parameters set to the method specified values? These include 400 µL sample volume, 480 second reaction time, 880 nm wavelength, 130 µL molybdate, and 20 µL ascorbic acid.  After analysis is completed, is the cuvette washed with alkaline EDTA solution to remove reagent deposits?  Is ammonia-free ASTM Type II water or better used for all solutions?	Relevant Aspect of Standards		Υ	N	N/A	Comments	
values? These include 400 µL sample volume, 480 second reaction time, 880 nm wavelength, 130 µL molybdate, and 20 µL ascorbic acid.  After analysis is completed, is the cuvette washed with alkaline EDTA solution to remove reagent deposits?  Is ammonia-free ASTM Type II water or better used for all solutions?	monitoring programs, are samples NOT filtered, but cooled to 4°C and analyzed within 48 hours? (No acid is	8.5					
alkaline EDTA solution to remove reagent deposits?  Is ammonia-free ASTM Type II water or better used for all solutions?  7.1	values? These include 400 µL sample volume, 480 second reaction time, 880 nm wavelength, 130 µL	17.1					
solutions?		11.2					
Notes/Comments:		7.1					